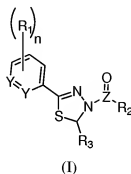


Amendments to the Claims

The listing of claims will replace all prior versions, and listings of claims in the application:

Listing of claims:

Claim 1. (Currently amended) A compound of Formula I:



in which

n is selected from 1, 2 and 3;

Z is selected from C and S(O); each

Y is independently selected from $-\text{CR}_4=$;

wherein R_4 is selected from hydrogen, cyano, hydroxyl, C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl and halo-substituted- C_{1-6} alkoxy;

R_1 is selected from halo, cyano, hydroxyl, C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl, halo-substituted- C_{1-6} alkoxy and $-\text{C}(\text{O})\text{OR}_4$; wherein R_4 is selected from hydrogen, cyano, hydroxyl, C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl and halo-substituted- C_{1-6} alkoxy;

R_2 is selected from C_{6-10} aryl, and C_{3-12} cycloalkyl; wherein any aryl or cycloalkyl of R_2 is optionally substituted with 1 to 5 radicals independently selected from halo, hydroxy, cyano, nitro, C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl, halo-substituted- C_{1-6} alkoxy, $-\text{C}(\text{O})\text{NR}_5\text{R}_5$, $-\text{OR}_5$, $-\text{OC}(\text{O})\text{R}_5$, $-\text{NR}_5\text{R}_6$, $-\text{C}(\text{O})\text{R}_5$ and $-\text{NR}_5\text{C}(\text{O})\text{R}_5$;

wherein:

R_5 and R_6 are independently selected from hydrogen, C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl, halo-substituted- C_{1-6} alkoxy, C_6 -

C_{10} aryl- C_{0-4} alkyl, and C_{3-12} cycloalkyl- C_{0-4} alkyl; wherein any aryl or cycloalkyl of R_5 is optionally substituted with 1 to 4 radicals independently selected from halo, hydroxy, cyano, nitro, C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl and halo-substituted- C_{1-6} alkoxy; R_3 is selected from C_{6-10} aryl and C_{3-12} cycloalkyl; wherein any aryl or cycloalkyl of R_3 is substituted with 1 to 5 radicals independently selected from halo, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkyl, halo-substituted- C_{1-6} alkoxy, -OXR₇, -OXC(O)NR₇R₈, -OXC(O)NR₇XC(O)OR₈, -OXC(O)NR₇XOR₈, -OXC(O)NR₇XNR₇R₈, -OXC(O)NR₇XS(O)₀₋₂R₈, -OXC(O)NR₇XNR₇C(O)R₈, -OXC(O)NR₇XC(O)XC(O)OR₈, -OXC(O)NR₇R₉, -OXC(O)OR₇, -OXOR₇, -OXR₉, -XR₉, -OXC(O)R₉, -OXS(O)₀₋₂R₉ and -OXC(O)NR₇CR₇[C(O)R₈]₂;

wherein:

X is selected from a bond and C_{1-6} alkylene wherein any methylene of X can optionally be replaced with a divalent radical selected from C(O), NR₇, S(O)₂ and O;

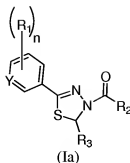
R_7 and R_8 are independently selected from hydrogen, cyano, C_{1-6} alkyl, halo-substituted- C_{1-6} alkyl, C_{2-6} alkenyl and C_{3-12} cycloalkyl- C_{0-4} alkyl;

R_9 is selected from C_{6-10} aryl- C_{0-4} alkyl and C_{3-12} cycloalkyl- C_{0-4} alkyl; wherein any alkyl of R_9 can have a hydrogen replaced with -C(O)OR₁₀; and any aryl or cycloalkyl of R_9 is optionally substituted with 1 to 4 radicals independently selected from halo, C_{1-6} alkyl, C_{3-12} cycloalkyl, halo-substituted- C_{1-6} alkyl, C_{1-6} alkoxy, halo-substituted- C_{1-6} alkoxy, -XC(O)OR₁₀, -XC(O)R₁₀, -XC(O)NR₁₀R₁₀, -XS(O)₀₋₂NR₁₀R₁₀ and -XS(O)₀₋₂R₁₀;

wherein:

R_{10} is independently selected from hydrogen and C_{1-6} alkyl;
~~and the pharmaceutically acceptable salts, hydrates, solvates and isomers thereof~~
or a pharmaceutically acceptable salt or isomer thereof.

Claim 2. (Previously presented) The compound of claim 1 of Formula Ia:



in which

n is selected from 1, 2 and 3;

Y is selected from $-\text{CH}=\text{}$;

R_1 is selected from halo, $\text{C}_{1-6}\text{alkyl}$, and $-\text{C}(\text{O})\text{OR}_4$; wherein R_4 is selected from hydrogen and $\text{C}_{1-6}\text{alkyl}$;

R_2 is selected from $\text{C}_{6-10}\text{aryl}$ and $\text{C}_{3-12}\text{cycloalkyl}$; wherein any aryl or cycloalkyl of R_2 is optionally substituted with 1 to 4 radicals independently selected from halo, hydroxy, $\text{C}_{1-6}\text{alkyl}$, halo-substituted- $\text{C}_{1-6}\text{alkyl}$ and $-\text{OC}(\text{O})\text{R}_5$; wherein R_5 is selected from hydrogen and $\text{C}_{1-6}\text{alkyl}$; and

R_3 is selected from $\text{C}_{6-10}\text{aryl}$ and $\text{C}_{3-12}\text{cycloalkyl}$; wherein any aryl or cycloalkyl of R_3 is substituted with 1 to 5 radicals independently selected from halo, hydroxyl, $\text{C}_{1-6}\text{alkoxy}$, halo-substituted- $\text{C}_{1-6}\text{alkyl}$, halo-substituted- $\text{C}_{1-6}\text{alkoxy}$, $-\text{OXR}_7$, $-\text{OXC}(\text{O})\text{NR}_7\text{R}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{XC}(\text{O})\text{OR}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{XOR}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{XNR}_7\text{R}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{XS}(\text{O})_{0-2}\text{R}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{XNR}_7\text{C}(\text{O})\text{R}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{XC}(\text{O})\text{XC}(\text{O})\text{OR}_8$, $-\text{OXC}(\text{O})\text{NR}_7\text{R}_9$, $-\text{OXC}(\text{O})\text{OR}_7$, $-\text{OXOR}_7$, $-\text{OXR}_9$, $-\text{OXC}(\text{O})\text{R}_9$ and $-\text{OXC}(\text{O})\text{NR}_7\text{CR}_7[\text{C}(\text{O})\text{R}_8]_2$;

wherein

X is selected from a bond and $\text{C}_{1-6}\text{alkylene}$;

R_7 and R_8 are independently selected from hydrogen, cyano, $\text{C}_{1-6}\text{alkyl}$, halo-substituted- $\text{C}_{1-6}\text{alkyl}$, $\text{C}_{2-6}\text{alkenyl}$ and $\text{C}_{3-12}\text{cycloalkyl}-\text{C}_{0-4}\text{alkyl}$;

R_9 is selected from $\text{C}_{6-10}\text{aryl}-\text{C}_{0-4}\text{alkyl}$ and $\text{C}_{3-12}\text{cycloalkyl}-\text{C}_{0-4}\text{alkyl}$;

wherein any alkyl of R_9 can have a hydrogen replaced with

$-\text{C}(\text{O})\text{OR}_{10}$; and any aryl or cycloalkyl of R_9 is optionally substituted

with 1 to 4 radicals independently selected from halo, C₁₋₆alkyl, C₃₋₁₂cycloalkyl, halo-substituted-C₁₋₆alkyl, C₁₋₆alkoxy, halo-substituted-C₁₋₆alkoxy, -XC(O)OR₁₀, -XC(O)R₁₀, -CR₁₀(NR₁₀R₁₀)=NOR₁₀, -XC(O)NR₁₀R₁₀, -XS(O)₀₋₂NR₁₀R₁₀ and -XS(O)₀₋₂R₁₀;

wherein

R₁₀ is independently selected from hydrogen and C₁₋₆alkyl.

Claim 3. (Previously presented) The compound of claim 2 in which

R₁ is selected from fluoro, chloro, methyl and -C(O)OCH₃; and

R₂ is selected from phenyl, cyclohexyl, cyclopentyl, and naphthyl; wherein any aryl or cycloalkyl of R₂ is optionally substituted with 1 to 4 radicals independently selected from fluoro, chloro, bromo, hydroxy, methyl, ethyl, propyl, t-butyl, amino, dimethyl-amino, methoxy, trifluoromethyl, trifluoromethoxy and -OC(O)CH₃.

Claim 4. (Previously presented) The compound of claim 3 in which R₃ is phenyl substituted with 1 to 5 radicals independently selected from fluoro, chloro, bromo, methoxy, hydroxyl,

difluoromethoxy, -OCH₂C(O)NH₂, -OCH₂C(O)OCH₃, -OCH₂C(O)NHCH₃, -OCH₂C(O)N(CH₃)₂, -R₉, -OR₉, -OCH₂R₉, -OCH₂C(O)R₉, -OCH₂C(O)NHR₉, -OCH₂C(O)N(CH₃)R₉, -OCH₂C(O)NHCH₂R₉, -OCH₂CN, -OCH₂C₂H₅, -OCH₂C₂H₄, -O(CH₂)₂OH, -OCH₂C(O)NH(CH₂)₂C(O)OC₂H₅, -OCH₂C(O)NH(CH₂)₂CH₂F, -OCH₂C(O)NHCH₂CH₂F, -OCH₂C(O)NH(CH₂)₂C(O)OH, -OCH₂C(O)NHCH(CH₂R₉)C(O)OC₂H₅, -OCH₂C(O)NHC(O)(CH₂)₂C(O)OCH₃, -OCH₂C(O)NH(CH₂)₂NHC(O)CH₃, -OCH₂C(O)NHCH₂C(O)C₂H₅, -OCH₂C(O)NH(CH₂)₂C(O)OC₄H₉, -OCH₂C(O)NHCH₂C(O)OC₂H₅, -OCH₂C(O)NHCH[C(O)OC₂H₅]₂, -S(O)₂CH₃, -OCH₂C(O)NHCH₂CF₃, -OCH₂C(O)NHCH₂C(O)(CH₂)₂C(O)OCH₃, -OCH₂C(O)N(CH₃)CH₂C(O)OCH₃, -OCH₂C(O)NH(CH₂)₃OC₂H₅, -OCH₂C(O)NH(CH₂)₃OCH(CH₃)₂, -OCH₂C(O)NH(CH₂)₂SCH₃, -OCH₂C(O)NHCH₂CH(CH₃)₂, -OCH₂C(O)NHCH(CH₃)CH₂OH, -OCH₂C(O)NHCH₂CH(CH₃)C₂H₅, -OCH₂C(O)NHCH(CH₃)C(O)OC₂H₅, -OCH₂C(O)NHCH₂CH(CH₃)₂ and -OCH₂C(O)(CH₂)₃OCH(CH₃)₂;

wherein

R₉ is phenyl, cyclopropyl-methyl, phenethyl; wherein any alkyl of R₉ can have a hydrogen replaced with -C(O)OC₂H₅; wherein any aryl of R₉ is optionally substituted with 1 to 4 radicals independently selected from methyl, ethyl, cyclopropyl, methoxy, trifluoromethyl, -OC(O)CH₃, -COOH, -S(O)₂NH₂, -CH(NH₂)=NOH, -C(O)OC₂H₅, -CH₂C(O)OH, -CH₂C(O)OC₂H₅, -CH₂C(O)OCH₃, -C(O)OCH₃, -C(O)NH₂, -C(O)NHCH₃ and -C(O)CH₃.

Claim 5. (Original) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 in combination with a pharmaceutically acceptable excipient.

Claim 6. (Cancelled) ~~A method for treating a disease or disorder in an animal in which modulation of LXR activity can prevent, inhibit or ameliorate the pathology and/or symptomatology of the disease, which method comprises administering to the animal a therapeutically effective amount of a compound of Claim 1.~~

Claim 7. (Cancelled) ~~The method of claim 6 wherein the diseases or disorder are selected from cardiovascular disease, diabetes, neurodegenerative diseases and inflammation.~~

Claim 8. (Cancelled).

Claim 9. (Cancelled) ~~A method for treating a disease or disorder in an animal in which modulation of LXR activity can prevent, inhibit or ameliorate the pathology and/or symptomatology of the disease, which method comprises administering to the animal a therapeutically effective amount of a compound of Claim 1.~~

Claim 10. (Cancelled) ~~The method of claim 9 further comprising administering a therapeutically effective amount of a compound of Claim 1 in combination with another therapeutically relevant agent.~~

Claim 11. (Currently amended) The compound of claim I selected from:

